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Nat Suppose Foreyo Co. Ud.

CONJ-01/08 CONJ-01/48 CO90-03/00 (27-12-71).. PIGMENT DISPERSIONS - MADE BY POLYMERISING COPOLYMERISARIE MONOMERS WITH DYE AND STABILISER ...

Full Priority: 22-12-70; 23-8-71; (4)/JA 115351; 64177; 64178, 64179; 64180.

NEW

Finely divided pigment dispersions are made by polymerising copolymerisable unsatd, monomers (!) in an ore, liqued which dissolves the monomers but not the polymer, in the presence of a stabiliser which is compatible with or chemically bonded to the copolymer and which contains a long hydrocarbon group so that the product can be dispers ed in low polar or non polar org. liquids, to give a dispersion with a particle size of 0.1-5 μ . The dispersion is coloured with a dye, pref. a water soluble dye or a dye con plex contg. a material with an electrical charge opposite to that of the dye.

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G2=A4.

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SE, ADVANTAGES

Products are stable pigment dispersions useful for colperine plastics such as PVC and printing links, csp. for the produ, of fluorescent colours. Dispersions have excellent light stability.

DETAILS

Dispersing agent may be non-polar aliphatic or aromatic hydrocarbons such as mineral spirit, kerosine, petrol-eum naphtha, hexane, cyclohexane, etc. The stabilisor may be cpds, with an ester grp. and a more than bC alkyl, such as hexyl acrylate, dihexyl lumarate, dioetwlitaeonate, etc. or polymers of these; poly-inyl esters, liquid polybutadiene The unsaid, monomers (I) ma urcthane -treated oils may be (A! those with low affinity for dyes such as npropyl(meth)acrylate, styrene, (B) copolymerisate mon-omers with 1 or 2 CN grps, such as vinyling e evanide. with high affinity to dyes; or (C) monomers with a tunction! al grp. having high affinity for dyes, such as methallyl sulphonic acid, (meth) acrylic acid, sinyl pyridine, diacetoneacrylamide, etc. (basic grps.). The heat and chemical resistance of the copolymer can be improved by the

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inclusion of monomer (D) such as diallyl phthalate, divinyl as 2,6-di tert, butyl 4-methylphenol and UV absorbers benzene, pentaerythritol tetramethacrylate, etc. Preferr ed copolymers contain 95/5 to 20/80 A:B, 99/1 to 70/30 A:C, or 90-20/80-5/30-1 A:B:C, opt. with 0.1-5 wt.% D (on A+B+C). Generally the dispersion contains 15-60 wt.% (esp. 25-60 wt.%) A+B+C. Polymen, is initiated with e.g.

benzoyl peroxide, AZBN. A wide range of dyes may be used, e.g. amino grp. -contg. dyes with cationic properties in dil acid soln, water sol, dyes contg. sulphonic acid or carboxylic acid grps.; water sol, dyes which are absorbed directly into fibres, water (in)soluble clear fluorescent dves and dves contg, no water-soluble grps, and which are soluble in oils and fats, hydrocarbons and molar solvents, mainly of the azo type but also some anthraquinonenigrosine - and azine types. The dyes are used at 0.3-25 wt. on copolymerisable monomers. Dye complexes can be made, esp. by using anionic surface active agents with cationic dyes and brightening agents, or a cationic surface active agent with acid dves and anionic fluorescent brightening agents. 1-15% of a HCHO contg. polycondensate which is sot, in monomers of type It but not in the org. liquid, may be added to improve the solubility and dispersibility of the pigment dispersions. The light resistance of the products can be increased by adding antioxidants such

such as phenyl salicylate. Polymsn, is carried out at 50-120°C (60-90°C) for 3-20 hrs. (8-12 hrs.).

E XA MPLE

A soln. of 459 pts, mineral spirit, 200 pts, sova hean oil modified alkyd resin (oil length 70), 3,5 pts. "Aizene Catilon Orange RH" and 2.0 pts. benzoyl peroxide was stirred at 80°C and a mixture of 200 pts. methyl arrylate and 150 pts, acrylonitrile added, over 4 hrs, Polymen, was continued for 6 hrs. to give an orange pigment dispersion.

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